THE INTERNATIONAL NETWORK ON RARE SKIN DISEASES
FOR PROFESSIONALS AND PATIENTS

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OUR MISSION

Since 2003, the Rare Skin Diseases Network of the Fondation René Touraine gathers dermatologists, scientists, patient representatives, scientific societies and public health authorities. Together, they develop new strategies to improve healthcare and social support for patients with rare skin diseases.

OUR ACHIEVEMENTS FOR 15 YEARS

• More than 15 conferences on diagnosis and management organized in European, Mediterranean and Middle-Eastern countries
  • More than 1000 healthcare professionals trained
  • More than 50 fellowships
• An international directory for patients and healthcare professionals
• An international Art Contest to raise awareness on rare skin diseases
• Launch of the ERN-Skin and support to the ERN-Skin coordination team

OUR OBJECTIVES FOR 2020

<table>
<thead>
<tr>
<th>Capacity Building</th>
<th>Training</th>
<th>Communication</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pilot centre for Epidermolysis Bullosa patients in Syria (questioned due to Covid-19)</td>
<td>On-site training in expert centers in France and Italy</td>
<td>Art Contest</td>
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<td>Pilot research project on phenotype-genotype correlation in Kuwait, Lebanon, Syria and Tunisia</td>
<td>Regional rare skin disorders symposium in Tunisia and Iran</td>
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</tbody>
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